

AMENDMENTS TO THE CLAIMS

By the present amendment, claims 30 and 40 are amended and claims 47-51 are added. This listing of claims will replace all prior versions and listings of claims in the application:

1-28. (Cancelled)

29. (Previously Amended) A method of producing L-β-lysine, comprising:

(a) culturing a prokaryotic host cell comprising an expression vector that encodes lysine 2,3-aminomutase in the presence of L-lysine, wherein the cultured host cell expresses lysine 2,3-aminomutase, and

(b) isolating L-β-lysine from the cultured host cells.

30. (Currently Amended) A method of producing L-β-lysine, comprising:

(a) incubating L-lysine in a solution containing substantially pure~~purified~~ lysine 2,3-aminomutase, said solution containing all cofactors required for lysine 2,3-aminomutase activity; and

(b) isolating L-β-lysine from the incubation solution.

31. (Previously Amended) The method of claim 30, wherein the lysine 2,3-aminomutase has an amino acid sequence selected from the group consisting of (i) SEQ ID NO: 4, and (ii) a conservative amino acid variant of SEQ ID NO: 4.

32. (Original) The method of claim 31, wherein step (b) further comprises isolating L-β-lysine from L-lysine via chromatography.

33-35. (Cancelled)

36. (Previously Amended) The method of claim 29 wherein the vector that encodes lysine 2,3-aminomutase has a nucleic acid sequence of SEQ ID NO: 3.

37. (Previously Amended) The method of claim 29 wherein the isolated L- β -lysine is enantiomerically pure.

38. (Previously Amended) The method of claim 30 wherein the isolated L- β -lysine is enantiomerically pure.

39. (Previously Added) The method of claim 30 wherein the cofactors required for lysine 2,3-aminomutase activity comprise:

- (i) at least one of ferrous sulfate or ferric ammonium sulfate;
- (ii) pyridoxal phosphate;
- (iii) at least one of dehydrolipoic acid, glutathione or dithiothreitol;
- (iv) S-adenosylmethionine; and
- (v) sodium dithionite.

40. (Currently Amended) A method of producing L- β -lysine, comprising:
(a) immobilizing substantially pure lysine 2,3-aminomutase on a suitable support;
(b) activating the lysine 2,3-aminomutase with cofactors required for lysine 2,3-aminomutase activity; and

(c) contacting L-lysine with the immobilized lysine 2,3-aminomutase to produce L- β -lysine.

41. (Previously Amended) The method of claim 40 wherein the L-lysine is contacted with the immobilized lysine 2,3-aminomutase for a sufficient amount of time to produce enantiomerically pure L- β -lysine.

42. (Previously Added) The method of claim 37 further comprising separating the L- β -lysine from the L-lysine.

43. (Previously Added) The method of claim 42 wherein the separation of the L- β -lysine from the L-lysine is achieved using high performance chromatography.

44. (Previously Added) The method of claim 37 wherein the process is a continuous process.

45. (Previously Added) The method of claim 37 wherein the cofactors required for lysine 2,3-aminomutase activity comprise:

(i) at least one of ferrous sulfate or ferric ammonium sulfate;

(ii) pyridoxal phosphate;

(iii) at least one of dehydrolipoic acid, glutathione or dithiothreitol;

(iv) S-adenosylmethionine; and

(v) sodium dithionite.

46. (Previously Amended) The method of claim 37, wherein the lysine 2,3-aminomutase has an amino acid sequence selected from the group consisting of (i) SEQ ID NO: 4 and (ii) a conservative amino acid variant of SEQ ID NO: 4.

47. (New) A method of producing L- β -lysine, comprising:

(a) incubating L-lysine in a solution containing purified lysine 2,3-aminomutase, wherein the lysine 2,3-aminomutase has an amino acid sequence selected from the group consisting of (i) SEQ ID NO: 4, and (ii) a conservative amino acid variant of SEQ ID NO: 4, said solution containing all cofactors required for lysine 2,3-aminomutase activity; and

(b) isolating L- β -lysine from the incubation solution.

48. (New) The method of claim 47, wherein step (b) further comprises isolating L- β -lysine from L-lysine via chromatography.

49. (New) A method of producing L- β -lysine, comprising:

(a) incubating L-lysine in a solution containing purified lysine 2,3-aminomutase other than that from *Clostridium subterminale* SB4, said solution containing all cofactors required for lysine 2,3-aminomutase activity; and

(b) isolating L- β -lysine from the incubation solution.

50. (New) The method of claim 49 wherein the isolated L- β -lysine is enantiomerically pure.

51. (New) The method of claim 49 wherein the cofactors required for lysine 2,3-aminomutase activity comprise:

- (i) at least one of ferrous sulfate or ferric ammonium sulfate;
- (ii) pyridoxal phosphate;
- (iii) at least one of dehydrolipoic acid, glutathione or dithiothreitol;
- (iv) S-adenosylmethionine; and
- (v) sodium dithionite.